

D'YAKONOV, I.A.; GOLCDNIKOV, G.V.; REPINSKAYA, I.E.

Reaction of diazoacetic ester with trimethylvinylsilane
catalyzed by copper sulfate. Zhur.ob.khim. 32 no.10:3450-3451
O '62. (MIRA 15:11)

1. Leningradskiy gosudarstvennyy universitet.
(Serine) (Silane)

GEORGEVICH, V. I.

MA Biological Institute for the Study of the Pathology of the Human Body, 1961.
Inst of Agriculture, Acad Sci USSR, Tashkent, 1961. 100 p., 100 p.

100 p., 100 p.

BORISOVA, A.G.; BOCHANTSEV, V.P.; BUTKOV, A.Ya., dotsent; VASIL'KOVSKAYA, A.P.;
VVEDENSKIY, A.I., dotsent; GOLODKOVSKIY, V.L.; GONCHAROV, N.F.
[deceased]; DROBOV, V.P., professor; KOROTKOVA, Ye.Ye.; KOSTINA, K.F.;
KUDRYASHEV, S.N. [deceased]; LAKHINA, M.M.; LINCHEVSKIY, I.A.;
MIRONOV, B.A. [deceased]; PAZIY, V.K.; POYARKOVA, A.I.; PROTOPOPOV,
G.F.; SUMNEVICH, G.P. [deceased]; KHAL'ZOVA, K.P.; YUZEPCHIK, S.V.;
KOROVIN, Ye.P., professor, glavnyy redaktor; ZAKIROV, K.Z., professor,
redaktor; SHIPUKHIN, A.Ya, redaktor izdatel'stva

[The flora of Uzbekistan] Flora Uzbekistana. Glav. red. Ye.P.Korovin.
Tashkent, Izd-vo Akademii nauk UzSSR. Vol.3. 1955. 825 p. (MLHA 9:10)

1. Deystvitel'nyy chlen AN UzSSR (for Korovin)
(Uzbekistan---Botany)

GOLODKOVSKIY, V.L.; AZIMOV, Kh.U.

Determining irrigation requirements of alfalfa grown for seed
production. Dokl. AN Uz. SSR no. 9: 39-41 '56. (MIRA 12:6)

1. Institut sel'skogo khozyaystva AN UzSSR. Predstavleno chlenom-
korrespondentom AN UzSSR S.S. Sadykovym.
(Alfalfa--Water requirements)
(Seed production)

GOLODKOVSKIY, V.L.; MELENT'YEVA, Ye.V.

Dynamics of the destruction of alfalfa plants by cuttings during the first year of life as related to varying densities of stand and irrigation conditions. Izv. AN Uz. SSR no. 10:45-50 '56.

(MIRA 14:5)

(Alfalfa)

GOLODKOVSKIY, V.L.; SHAAKRAMOV, K.Sh.

Development of aerial parts and the root system in perennial
grasses. Uzb. biol. zhur. no.1:36-41 '61. (MIFA 14.3)

1. Institut genetiki i fiziologii rasteniy AN UzSSR.
(GRASSES)

GOLODKOVSKIY, V.L.; POPOVA, Ye.A.

In memory of V.S.Shardakov. Uzb. biol. zhur. no.1:77-78 '61.
(MIRA 14:3)

(SHARDAKOV, VASILII SEMENOVICH, 1901-1960)

GOLOVCHENKO, S.G.; GOLODKOVSKIY, V.L., kand.sel'khoz.nauk, otv. red.;
GOR'KOVYI, P.I., red.; KARABAYEVA, Kh.U., tekhn. red.

[Feed production in piedmont areas; research results] Kormodoby-
vanie v predgor'iax; opyt issledovaniia. Tashkent, Izd-vo Akad.
nauk UzSSR, 1962. 166 p. (MIRA 15:7)
(Uzbekistan--Forage plants)

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GOLEKOVSKIY, V.I.; PATEBNIATOV, S.I.

Heterosis in crossing self-pollinated lines of corn. Dok.
biol. zhur. 9 no.1:65-66 '65. (ENLA 18:c)

I. Institut eksperimental'noy biologii tekhnicheskikh nauchnykh
kul'tur AN USSR.

PROF. M. I. L. ... L. A., COLOMBIA, S. I.

Investigation of the ... of the acoustical
material based on polyvinyl chloride resin. First, massy
not ... (MIRA 1974)

1. NAME, I.M.; 2. BIRTHDAY, I.M.; 3. BIRTHDAY, I.M.; 4. BIRTHDAY, I.M.;
5. BIRTHDAY, I.M.

Removal of the subject from the unit with a view to the
of certain of the subject's activities.

1944

ISPOLITOV Yu. (author) ... (M. L. V. ...), Engineer
Podolsk 11

New scientific ... from the Ilkhachov Platform.
Tel ... no 4 ... (M. L. V.)
... (instruction)

GALONIKOV, A. V.

1. Univ. of Kiev - 1951-52

1951-52

General Scientific Section - 1951-52, in 1951, A. V. Galonikov,
T. A. Galonikova, A. V. Galonikov, A. V. Galonikov, and A. V. Galonikov

West Lenin and U, 1951-52, Vol. 1, No. 2, pp 198-199

The General Scientific Section of Leningrad University was, from 1951-52, 1953-54.
The Math. section was subdivided into math, mechanics, and astronomy; the physics
comprised also applied physics. The chemistry section dealt also with cooperation with
industry.

1951-52

GOLODNIKOV, G.V.

USSR/Chemistry - Catalysis

Card 1/1 Pub. 151 - 12/33

Authors : Dolgov, B. N., and Golodnikov, G. V.

Title : Catalytic derivation of mixed ketones from primary alcohols. Part 1.-
Ketonization of an ethyl and n-butyl alcohol mixture

Periodical : Zhur. ob. khim. 24/6, 987-993, June 1954

Abstract : Experiments on the ketonization of an ethyl and n-butyl alcohol mixture, carried out in the presence of additionally introduced hydrogen in molar ratio of 1 : 1 to the alcohol mixture, are described. The effect of the hydrogen addition on the ketone yield and catalyst activity is explained. The conditions favorable for such ketonization process and the experimental results obtained are listed. Nine references: 6-USSR and 3-USA (1924-1951). Tables.

Institution : The A. A. Zhdanov State University, Leningrad

Submitted : December 26, 1953

GOLDNIKOV, G. V.

3
CII ✓ Catalytic preparation of mixed ketones from primary
alcohols. II. Ketonization of a mixture of ethyl and iso-
amyl alcohols. R. N. Dulgoy and G. V. Goldnikov.
J. Gen. Chem. U.S.S.R. 24, 1169-61 (1954) (Engl. transla-
tion).—See C.A. 49, 12282a. B. M. R. ①

64-1-1-1

USSR .

62
D
/ Catalytic preparation of mixed ketones from primary
alcohols. II. Ketinization of a mixture of ethyl and isocamyl
alcohols. B. N. Dolgov and G. V. Golodnikov (Leningrad
State Univ.). *Zhur. Obshchei Khim.* 24, 1167-70 (1954);
cf. C.A. 49, 8794f. —Passage of EtOH and iso-AmOH in
2:1 molar ratio over Cu catalyst activated according to
directions given by D. and Bolotov (*Vestnik Leningrad.
Gosudarst. Univ.* S. No. 6 (1950) at 320° and space velocity
50-150 gives the best yield (18-21%) of iso-BuAc, with 0.6-
1% Me₂CO and about 5% (iso-Bu)₂CO. At 275-300°
are formed esters in yields near 45%; these include EtOAc,
iso-AmOAc, iso-BuCO₂Et and iso-BuCO₂CH₂CH₂CH₂Me.
The effluent gases from the ketonization contain mostly H₂,
with small amounts of CO, CO₂ and olefins. G. M. K.

G. V. Golodnikov

USSR

Catalytic preparation of mixed ketones from primary alcohols. III. The ketonization of a mixture of butyl and isoamyl alcohols. B. N. Dolgov and G. V. Golodnikov (Leningrad State Univ.). *Zh. Priklad. Khim.* 24, 1461-71 (1954); cf. *Chem. Abstr.* 49, 5:94f.---In the passage of mixed BuOH and iso-AmOH over activated Cu catalyst at 275-425°, the best yields (15%) of $\text{PrCOCH}_2\text{CHMe}_2$, 23% Pr_2CO , and 2-3% iso-Bu₂CO are obtained when the molar ratio of BuOH to iso-AmOH is 2-3:1 and the temp. is 350-75°, with a space velocity of 100-50. At lower temps. (275-335°) the main products are esters forming up to 60% of total product. The off-gas consists mainly of H₂ with small amts. of CO, CO₂, and olefins. The results indicate that aldehydes and esters are the intermediate products of ketonization. The correct mechanism appears to be that suggested by Sabatier-Kagan (no ref. cited), in which esters are intermediates. The reaction yields aldehydes, which condense to esters which are then decarboxylated with elimination also of ROH and olefins, along with the H₂ formed in the initial dehydrogenation of ROH. G. M. Kozlovskii...

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GOLODNIKOV, G. V.

7
 ✓ Decomposition of esters of benzoic acid over chromium catalyst. B. M. Dolgov, G. V. Golodnikov, and L. M. Chernykhina (State Univ. Leningrad, USSR, *Dokl. Akad. Nauk SSSR*, 1955, 60 (1955).—Passage of $BzOPh$, $BzOBu$, and $BzOAm$ over $Zn-Cr$ catalyst (cf. C.I. 40, 10170c) at 380–420° yields C_6H_6 , CO_2 , and olefin, in one reaction sequence, and an alkyl aryl ketone in another sequence. For prepn. of the ketones the best conditions are 400–20° 31. square velocity 50 (9); under these conditions the yields of ketones are: $AcPh$ 0.3%, $PhCOPh$ 6.6%, $i-BuCOPh$ 3.3%. G. M. Kosolapoff

USSR/Organic Chemistry. Synthetic Organic Chemistry.

E-2

Abs Jour: Ref Zhur-Khimiya, No 6, 1957, 19254

Author : Dolgov N. N., Golodnikov G. V., Glushkova N. Ye.

Inst :

Title : Conversion of Tetraethylsilane over a Chromium Catalyst.

Orig Pub: Zh. obschch. khimiyi, 1956, 26, No 6, 1688-1691

Abstract: At 530-630° tetraethylsilane (I) over a chromium catalyst
in an atmosphere of H_2 is decomposed into triethylsilane
and C_2H_4 . At 600-630° a deep hydrogenolysis ensues with
the formation of C_2H_5 and Si. Dehydration of I to triethyl-
vinylsilane is not observed.

Card : 1/1

GOLODNIKOV, G. V.

✓ Transformation of biological data into a form suitable for
B. S. Datsar, B. S. Golodnikov, B. S. Golodnikov, B. S. Golodnikov
J. Gen. Chem. 1955, 26, 1331-5 (1955) (1955) (1955) (1955)
Lithuanian SSR, 1955

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PHASE I BOOK EXPLOITATION

614

~~Golodnikov, Gennadiy Vladimirovich, Nizovkina, Tat'yana Vsevolodovna
and Ryskal'chuk, Apollinariya Terent'yevna~~

Praktikum po organicheskomu sintezu (Practical Work in Organic Synthesis) Leningrad, izd-vo Leningrad. univ-ta, 1957. 137 p.
6,080 copies printed.

Sponsoring Agency: Leningrad. Universitet.

Ed. (title page): Dolgov, B.N., Professor; Ed. (inside book):
Shchemeleva, Ye.V.; Tech. Ed.: Vodolagina, S.D.

PURPOSE: This manual is intended for the use of chemistry students taking a laboratory course in organic synthesis.

COVERAGE: This manual is presented as a guide to practical laboratory work in organic synthesis. It is divided into three parts. The first part is devoted to general methods and procedures of laboratory work in this field. Particular attention is given to

Card 1/E

Practical Work in Organic Synthesis

014

fractional distillation, fractionating columns, steam distillation, reduced-pressure distillation, recrystallization, and the determination of melting points. Instructions are given for keeping records, safeguarding against accidents, what to do in case of fire, first aid in case of burns, glass cuts, etc. The second part describes the synthesis of organic compounds. The syntheses are divided according to types of reaction. For each type of reaction a few examples are given distinguished one from the other according to methods of separation and treatment of organic compounds produced. The third part of the manual includes supplementary syntheses which are given to laboratory students as finals. Every synthesis described in the manual has been checked and proved many times over during many years by the faculty of organic chemistry at the Leningrad State University. There are no personalities and no references.

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GOLODNIKOV, G. V.

Caustic preparation of leucine from a substance of α -amino acid and α -hydroxy acid. E. N. Doherty, G. V. Lusk, *Alkyl and Ye. A. Forster*, *Chem. Ind. (London)*, 1949, 1308. Observed m.p. 170-171°C. (decolor.) Fusion of α -OH. Boil. over CaH_2 carbide (of α -amino). Distribution. Lentin and Skell. *Anal.* 1949, 104 at 340-500° (melt up to 200°). MeCO Pr. distill. with 1% MeCO and 10% PrCO . Fraction scheme of the form. Reaction of products and finally discarded. Formation of MeCO Pr. probably caused by decomposition of AsH with PrCHO or MeCO . The reason for variation (temp. and reactant proportions) (A. I. and B. I.) are tabulated. G. M. Knechtel.

Distri 484

20-6-18/47

AUTHORS: Dolgov, B. N. , Golodnikov, G. V. , and Golodova, L. G.

TITLE: On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons (O vozmozhnosti kataliticheskogo dehidrirovaniya kremsko-levodorov)

PERIODICAL: Doklady AN SSSR, 1957, Vol. 117, Nr 6, pp. 987 - 989 (USSR)

ABSTRACT: There exist no references to this kind of dehydrogenation of silicon-paraffins in publications. In reference 1 it is reported that under the conditions selected there the above-mentioned reaction with tetraethylsilane did not take place. The authors succeeded in finding a catalyst (placed at their disposal by Yu. A. Gorin and S. M. Monoszon) and in determining the conditions of the dehydrogenation of a mixed tetra-alkylsilane, namely trimethylbutylsilane. The nature of the above-mentioned catalyst is not mentioned in the paper. At 550 - 575 °C 6,2 - 8,6 % yields of trimethyl-butetyl-silane, calculated on the trimethylbutylsilane sent through, were obtained (table 1). The catalyst is highly stable: neither the yields of silicon-olefin change nor is silicon deposited on the catalyst. The above-mentioned yields can still be increased by repeated passage of condensates over the catalyst, as the latter contain considerable quantities of unchanged trimethylbutylsilane.

Card 1/3

20-6-18/47

On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons

Beside the dehydrogenation, especially at high temperatures (575 - 600°C), some side reactions take place which are connected with the thermal decomposition of trimethylbutylsilane. Of special interest is the formation of tetramethylsilane and propylene which occurs under splitting up of the C-C bond in the butyl radical. At the same temperatures a destructive hydrogenation of the formed tetramethylsilane by hydrogen, produced in the dehydrogenation of trimethylbutylsilane takes place. Theoretically the following isomers of trimethylbutenyl-silane are possible: $(CH_3)_3SiCH=CHCH_2CH_3$ (cis- and trans-forms) (I), $(CH_3)_3SiCH_2CH=CHCH_3$ (cis- and trans-forms) (II), and $(CH_3)_3SiCH_2CH_2CH=CH_2$ (III). Of these, however, only trimethyl- γ -butenylsilane (III) is known. The authors did not succeed in isolating the silicon olefin in a pure state, as the boiling points of all products and of the initial substance are supposed to lie very close to each other. The constants of the fraction 109 - 111°C, most enriched with silicon-olefin, are in table 2 compared with the properties of the known γ -isomer (III) and of the initial substance. The silicon-olefin obtained by the authors apparently is the γ -isomer (I). The absence of the β -isomer (II) is confirmed by the speed of the reduction of the produced silicon-olefin. Finally the absence of the β -isomer is confirmed by the production of a stable dibromide

Card 2/3

20-6-18/47

On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons

of trimethyl-butenyl-silane. A kind of short experimental part with the usual data is given which is not designated as such. There are 2 tables, and 6 references, 5 of which are Soviet.

ASSOCIATION:

Leningrad State University ineni A. A. Zhdanov
(Leningradskiy gosudarstvennyy universitet in. A. A. Zhdanova)

PRESENTED: August 5, 1957, by A. V. Lopchikov, Academician

SUBMITTED: August 5, 1957

AVAILABLE: Library of Congress

Card 3/3

AUTHORS: Golodnikov, G. V., Delgov, B. N., SCV/70-28-6-21/66
Propp, L. N.

TITLE: Catalytic Synthesis of Ketones From a Mixture of n-Butyric
Acid and Ethyl Alcohol (Kataliticheskoye polucheniye ketonov
iz smesi n.-maslyanoy kisloty i etilovogo spirta)

PERIODICAL: Zhurnal obshchey khimii, 1958, Vol. 26, Nr 8,
pp. 2097 - 2099 (USSR)

ABSTRACT: The authors showed already in the previous paper (Ref 1)
that a methyl-propylketone and besides this symmetrical
ketones, i.e. acetone and dipropyl ketone, are formed in
in the case of passing of a mixture of acetic acid and n-butyl
alcohol over a Cr-Mn catalyst. The suggested reaction scheme
is based upon a dissociation of the primarily formed ester,
of butyl acetate into aldehydes, their aldol-condensation
and the further transformation of aldol into methyl-propyl
ketone. The present paper is a continuation of the experiments
which try to synthesize mixed ketones from mixtures of acids
and alcohols. A mixture of ethyl alcohol and n-butyric acid
was chosen. This mixture is interesting since the assumed

Card 1/3

Catalytic Synthesis of Ketones From a Mixture of
n-Butyric Acid and Ethyl Alcohol

SOV/77-28-8-21/66

intermediate product formed in the ketone formation, the ethyl butyrate is metameric to the butyl acetate which for its part represents an intermediate product which was formed in the ketone formation of the mixture of acetic acid n.-butyl alcohol. Thus the transformations of these esters which are in the metamerismic ratio are bound to lead to one and the same aldehydes, i.e. to the acetic- and butyric aldehyde. Therefore the formation of the same ketones, i.e. of methyl-propyl ketone, acetone, and dipropyl ketone observed in the reaction confirms to a certain extent the correctness of the suggested mechanism of the ketonization of the mixtures of acids and primary alcohols. The formation of a mixed ketone is demonstrated in the given reaction scheme, where attention must be paid to R and R' of both mixtures. The conditions and results of the experiments are given in tables 1,2,3, the properties of the ketones in table 4. Cr-Mn catalyst was used as a catalyst in this work. There are 4 tables and 4 references, which are Soviet.

Card 2/3

Asymmetric Synthesis of Ketones From a Mixture of
Acetic Acid and Ethyl Alcohol

SCV/79-28-6-21/66

ORIGIN: Leningradskiy gosudarstvennyy universitet (Leningrad State
University)

SUBMITTED: July 2, 1957

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53700 2209

AUTHORS: Dolgov, B. N. (Deceased), Tolstunov, G. V., Gensler, I. B.

TITLE: Catalytic Conversion of Tetraalkyl Silanes. III. Catalytic
Dehydrogenation of Trimethyl Propyl-, Triethyl Butyl-,
and Trimethyl Hexyl Silane

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 3,
pp. 2988-2995

TEXT: In continuation of their previous paper (Ref. 2), the authors studied the dehydrogenation of silicon hydrocarbons of the series $(CH_3)_3SiR$, ($R = C_3H_7, C_4H_9, C_6H_{13}$). The investigations were carried out with the catalyst used in the previous investigation (Ref. 2) by Yu. A. Gorin and S. M. Monozon and placed at the authors' disposal. The tetraethyl silane passed through over this catalyst at 500-600° (volume rate 30) was not dehydrogenated: Besides unchanged tetraethyl silane, triethyl silane and diethyl silane, that are formed by the splitting off of ethylene, were detected (Table 1). The gaseous products consisted of hydrogen, saturated hydrocarbons and ethylene. The silanes of the series

Card 1/3

Catalytic Conversion of Tetraalkyl Silanes 3/57/40, 830/001/006 018
III. Catalytic Dehydrogenation of Trimethyl 8001, 8064
Propyl-, Trimethyl Butyl-, and Trimethyl Hexyl Silane

$(\text{CH}_3)_3\text{SiR}$, however, contain no ethyl radicals and can therefore be dehydrogenated with the above catalyst. Table 2 shows the optimum conditions of the dehydrogenation of trimethyl butyl silane, and Table 3 those of the dehydrogenation of trimethyl propyl silane (depending on the temperatures and volume rates), as well as the yields of the reaction products. The reactions carried out at temperatures above 570°C in the presence of a catalyst proceeded according to Scheme 2 with the bonds C - C and C - Si undergoing cleavage. Only by multiple fractional distillation, in a rectifying column, it was possible to separate the fraction boiling between 84.0 and 84.5°C that (owing to the thiocyanogen number) contained 21.1% silicon olefin. The three intensive lines of the Raman spectrum of the fractions enriched with the silicon olefin are probably caused by the isomers $(\text{CH}_3)_3\text{SiCH}=\text{CHCH}_3$ and $(\text{CH}_3)_3\text{SiCH}_2\text{CH}=\text{CH}_2$. The dehydrogenation product of trimethyl propyl silane appears, according to its spectral analysis, as in the case of trimethyl butyl silane (Ref. 2), to consist for the major part of the isomer of silicon olefin that contains the double bond in

Card 2/3

Catalytic Conversion of Tetraalkyl Silanes 3/07/60/049/004/006/015
III. Catalytic Dehydrogenation of Trimethyl Propyl-, Trimethyl Butyl-, and Trimethyl Hexyl
Silane R001/R064

α -position to the silicon. A temperature of 540°C and the volume rate of 45 proved to be the optimum reaction conditions for the dehydrogenation of trimethyl hexyl silane (Table 4). Also here, the Raman spectrum shows in all cases the frequencies characteristic of the double bond. There are 8 tables and 17 references: 8 Soviet, 6 US, and 3 German.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet
(Leningrad State University) X

SUBMITTED: October 2, 1959

Card 3/3

54879

S/079/60/030/010/020/030
3001/3066

S 3700 4 1273, 1205

AUTHORS: Solodnikov, G. V. Dolgov, B. M. (Deceased) and
Sedova, V. F.

TITLE: Synthesis and Properties of Trialkyl-triaryl-(p-bromo-
phenoxy)-silanes. I. Trimethyl-, Triethyl- and Tri-
propyl-(p-bromo-phenoxy)-silanes

PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 10,
pp. 3352 - 3358

TEXT: Only the first member of silicon-containing bromides of the
general formula $p-R_2SiOC_6H_4Br$ ($R=CH_3$) has so far been described in pub-
lications (Ref.1). The authors of the present paper intended to syn-
thesize members of the same series with $R = C_2H_5, C_3H_7$, and to study
the possibilities of their synthesis, first of all, when using the
Grignard reagents (obtained from the above bromides) in the synthesis
of silicon-containing aromatic alcohols and acids. The hydrolytic sta-
bility of the ether group $Si-C-C_{ar}$ (C_{ar} = aromatic carbon of benzene)

Card 1/3

54579

Synthesis and Properties of Trialkyl-tri- S/C79/60/C33/C10/C2C/C3C
aryl-(p-bromo-phenoxy)-silanes. I Tri- B001/BC56
methyl-, Triethyl-, and Tripropyl-(p-bromo-phenoxy)-silane

should further be investigated. The trialkyl (p-bromo phenoxy)-silanes newly synthesized by the authors are stable to hydrolysis; the Si-O-C_{ar} bond remains unchanged (Ref. 2). The secondary alcohols were synthesized according to the Scheme $R_3SiOC_6H_4MgBr + CH_3CHO \rightarrow R_3SiOC_6H_4CH(OH)CH_3$. Alcohols with R = CH₃ or C₂H₅ however, were not obtained because of their hydrolytic instability. At R = CH₃, only hexamethyl disiloxane (30% yield) and a silicon-free resin resulted. When R was C₂H₅, triethyl-phenoxy-silane (24%), was found in addition to hexaethyl disiloxane (38%) and a large quantity of resin. Methyl-(p-triisopropyl-siloxy-phenyl) carbinol was far more stable (33%). Tripropyl-phenoxy silane (20%) and a small resin quantity were obtained as side products which indicates a comparative hydrolytic stability of the Si-O-C_{ar} bond in the molecule of the alcohol. Molecular weight and silicon content are in good agreement with the formula $(C_6H_4)_3SiOC_6H_4CH(OH)CH_3$. Further

Card 2/3

10-9

Synthesis and Properties of Trialkyl-arylsilanes. I. Tri-
 aryl-(p-bromophenoxy)silanes. I. Tri-
 methyl-, Triethyl-, and Tripropyl (p-bromophenoxy)silane

S/073/60/010/010/020/030

EC01/1066

experiments concerning the synthesis of the alcohols show that, when
 propyl radicals are introduced, the hydrolytic stability of Si-C_{ar}
 bonds increases so much that the corresponding alcohol can be separated.
 Acids of the general formula $\text{R}_2\text{Si}(\text{C}_6\text{H}_4\text{COOH})_2$ could not be synthesized
 by means of the stabilizing propyl radicals; p-oxybenzoic acid and
 silicon-containing cleavage products were found instead of these acids
 (Table 1). The effect of the magnitude of the radical on the silicon
 atom upon the properties of the Si-C_{ar} bond was experimentally deter-
 mined by cleavage of the Grignard reagents (Scheme 1). Table 1 illus-
 trates the cleavage of the Grignard reagents: $\text{R}_2\text{Si}(\text{C}_6\text{H}_4\text{MgBr})_2$. There are
 4 tables and 2 references, 6 Soviet, 3 US, and 1 German.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad
 State University)

SUBMITTED: December 2, 1959

Card 3/3

EL385

53700 2209, 1271, 1153

S/073/60/060/010/028/030
BCC/3066

AUTHORS: Golodnikov, G. V. and Reginskaya, I. B.
TITLE: Catalytic Dehydrocyclization of Trimethyl 1-hexyl Silane
PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 10,
pp. 3501 - 3502

TEXT: In order to find out whether dehydrocyclization of trimethyl hexyl silane is possible, the authors allowed it to pass over a chromium catalyst at 530-590°C (Ref.1). Under these conditions, dehydrocyclization gives trimethyl-phenyl silane:

$(CH_3)_3SiC_6H_{13} \rightarrow 4H_2 + (CH_3)_3SiC_6H_5$ The aromatization of silicon paraffins is a reaction hitherto unknown in organosilicon chemistry. An unsaturated hydrocarbon appears as intermediate, whose content in the condensates varied between 7.5 and 21%, according to the temperature (determined by the thiocyanation method). Trimethyl silane, tetramethyl silane, pentane, hexane and benzene were the side products. The condensates repeatedly underwent fractional distillation. The separated

Card 1/2

ELSES

Catalytic Dehydrocyclization of
Trimethyl-hexyl Silane

S/070/60/C30/010/028/030
B001/B066

fraction with a boiling point between 163 and 165° contained trimethyl-phenyl silane in addition to unchanged trimethyl hexyl silane. Ref. 2 gives the data published on trimethyl-hexyl silane, and Ref. 3 those on trimethyl-phenyl silane. The presence of trimethyl phenyl silane in the fraction boiling between 163 and 165° is confirmed by its refractive index and spectroscopic data. The infrared spectrum of the fraction shows an absorption maximum which is characteristic of the phenyl ring which is bound to the silicon (Ref.4) This study is being continued to find stronger catalysts for the dehydrocyclization of silicon paraffins. There are 4 references: 1 Soviet, 1 Japanese, and 2 US. ✓

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad
State University)

SUBMITTED: May 26, 1960

Card 2/2

S 01.61.031.01.001.001
0118.0101

AUTHORS

Golodnikov G. V. and Afanas'yev A. I.

TITLE

Tetramyl-, trihexyl- and triphenyl-(n-bromophenoxy)silanes

PERIODICAL

Zhurnal obshchey khimii 36: 30-31, 1966, 2335, 1 p.

TEXT

The authors studied the synthesis and properties of two little known silicoorganic bromides and two new silicoorganic alcohols: tetramyl-(n-bromophenoxy)silane (I); trihexyl-(n-bromophenoxy)silane (II); triphenyl-(n-bromophenoxy)silane (III); methyl-(n-bromophenoxyphenyl)carbinol - a $(C_5H_4)SiOC_6H_4CH(OH)CH_3$ (IV) and methyl-

(n-trihexylsiloxyphenyl)carbinol - a $(C_6H_{11})_3SiOC_6H_4CH(OH)CH_3$. In previous research by G. V. Golodnikov, B. N. Dolgov, A. F. Zolotarev, Zh. obshch. khimii 36: 3352, 1966) on the properties and condensation of bromides of the type $aR_3SiOC_6H_4Br$ disclosed the possibility of their

Card 1/3

Triamyl-, trihexyl- and

conversion into secondary silicoorganic alcohols by means of organochloride and organosilane compounds. The method of the synthesis of the condensation of trialkyl- and trialkylsilanes with organochloride and $ZnCl_2$ or $SnCl_4$ catalyst, given in N. A. Belyi, A. I. Anisimov, N. F. Khazantov (Ref. 12, AN SSSR, Otdel Khim. i Mekh., 1968) was used to prepare I, II and III. For syntheses of I and II, 10 g of Mg, abs. ether, I and ethyl bromide was freshly distilled, after which the solution was successively cooled, reacted with organochloride in abs. ether, boiled for a further hour, and the product was purified. The subsequent vacuum distillation of the product gave I, II and III, which yielded three fractions: 175-180°C, 180-190°C, and 190-200°C. IV contaminated with some organochloride and ethyl bromide. IV was converted into the secondary silicoorganic alcohols by the fraction boiling at 80-120°C, containing a small amount of organochloride and ethyl bromide, which was removed by distillation. The authors conclude that the organochloride and ethyl bromide are not alkylated by the organochloride and ethyl bromide.

Card 20

3-679-61-031 013 013 013
D22870407

Triamyl-, trihexyl-, and

Si-O-C bond in the molecules of IV and V, which may be due to the screening effect of the amyl and hexyl radicals linked with the Si atom. IV and V react with Na to give H_2 , and their IR (K) spectra contain absorption maxima at 3400 cm^{-1} and 3360 cm^{-1} respectively. There are 2 tables and 4 references: 2 Soviet-bloc and 2 non-Soviet bloc.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad State University)

SUBMITTED: December 6, 1960

Card 3/3

S/079/61/031/011/014/015
D228/D305

5 3700
AUTHORS:

Golodnikov, G. V., and Koroleva, G. N.

TITLE:

Catalytic dehydrogenation of trimethylethylsilane

PERIODICAL:

Zhurnal obshchey khimii, v. 31, no. 11, 1961, 3738-3740

TEXT: The authors studied the catalytic dehydrogenation of trimethylethylsilane—the first member of a group of mixed silanes with the formula Me_3SiR . B. N. Dolgov, G. V. Golodnikov, and I. B. Gensler (Ref. 1: Zh. obshch. khimii, 30, 2988, 1960), whose experimental procedure was followed in this work, also examined the dehydrogenation of other mixed silanes— Me_3SiPr , Me_3SiHx —and showed that tetraethylsilane, on the contrary, does not undergo dehydrogenation under the chosen conditions. The authors' data indicate that the reaction proceeds best at 590 – 600°, some 20 – 30° higher than is the case with silanes containing propyl, butyl, and hexyl radicals. Raising the temperature to 620° promotes the development of side-reactions—when trimethylsilane (I), trimethylvinylsilane (II),

Card 1/2

AUTHORS: Golodnikov, G. V. and Khalutina, A. A.

S:079/62/032/007/003/007
1032/1232

TITLE: Catalytic dehydrogenation of γ -trimethyl-silyl-propyl alcohol

PERIODICAL: Zhurnal obshchei khimii, v. 32, no. 7, 1962, 2302-2305

TEXT: The aim of this study was to work out a general method for the preparation of silicon-containing aldehydes by way of dehydrogenation of the corresponding primary alcohols. The best yield of β -trimethyl-silyl-propionic aldehyde was obtained by dehydrogenation of the corresponding alcohol over a catalyst designated "violet copper" (reduced copper) at 300 °C. The yield amounted to 76.7% of the alcohol that reacted, or to 37.8% of the total amount of alcohol. 45% of the alcohol did not react. Side reactions took place to a small extent. Dehydration gave trimethyl-allyl-silane (3.5%). Hexamethyl-disiloxane (2.3%) and propylene were also formed. There is 1 table. The English-language references read: 1) K. Frisch, and P. Shroff, J. Am. Chem. Soc., 75, 1249 (1953); Ch. A., 47, 9055 (1953). 3) C. Burkhard and D. Hurd, J. Org. Ch., 17, 1107 (1952). 4) C. Brannen Ch. A. 48, 624 (1954). 5) J. Speier, J. Webster, G. and Barnes, J. Am. Chem. Soc., 79, 574 (1957). 9) L. Sommer, R. Van Streen and F. Whitmore, J. Am. Chem. Soc., 71, 3056 (1949). 6) Beilste. II 2. 8. 11. 11. 11.

ASSOCIATION: Leningradskii gosudarstvennyi

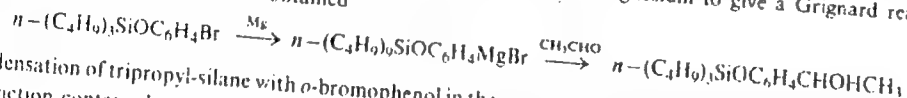
AUTHORS: Golodnikov, G. V. and Sycheva T. P.

S:079/62/032/007/004/007
I032/I232

TITLE: Synthesis and properties of tributyl-(*p*-bromophenyl)-silane

PERIODICAL: Zhurnal obshchei khimii, v. 32, no. 7, 1962, 2305-2307

TEXT: Tributyl-(*p*-bromophenoxy)-silane, a compound not described in the literature hitherto, was synthesized by catalytic condensation (involving dehydrogenation) of tributyl-silane with *p*-bromophenol, in the presence of SnCl_2 . Tributyl-(*p*-bromophenoxy)-silane reacted with magnesium to give a Grignard reagent, from which a carbinol could be obtained



Condensation of tripropyl-silane with *o*-bromophenol in the presence of SnCl_2 was attempted, but the products of reaction contained considerable amounts of hexapropyl-disiloxane, *o*-bromophenol, as well as 57.9% of the calculated amount of hydrogen. The tripropyl-(*o*-bromophenoxy)-silane is considered to have undergone hydrolytic dissociation.

ASSOCIATION: Leningradskii gosudarstvennyi universitet (Leningrad State University)

SUBMITTED: July 5, 1961

Card 1/1

GOLODNIKOV, G.V., D'YAKONOV, I.A.; REPINSKAYA, I.B.; FOMINA, O.S.

Copper sulfate catalyzed reaction of diazoacetic ester with
3-trimethylsilyl-1-propene and 4-trimethylsilyl-1-butene.

Zhur.ob.khim. 33 no.7:2422-2423 J1 '63.

(MIRA 16:3)

1. Leningradskiy gosudarstvennyy universitet.
(Silicon organic compounds) (Acetic acid)

GOLDENKOV, G.V.; SHAROVA, N.A., V.V.

Catalytic dehydrogenation of $\sqrt{\text{V}}$ -trialkylsilylpropyl alcohols.
Part 2. Zhur.ob.khim. 33 no.10:3262-3264, 0 '63.(MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.

GOLODNIKOV, G.V.; MARSH, G.A.

Synthesis and properties of trialkyl(p-bromophenoxy)silanes. Part 4.
Zhur.ob.khim. 33 no.10:3265-3266 O '63. (MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.

D'YAKONOV, I.A.; GOLODNIKOV, G.V.; REPINSKAYA, I.B.; FOMINA, O.S.

Reactions of diphenylmethylen and carbethoxycarbene with
alkenylsilanes. Zhur.ob.khim. 33 no.10:3436-3439 O '63.
(MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.

1. The first part of the document is a list of the names of the persons who were present at the meeting.

2. The second part of the document is a list of the names of the persons who were present at the meeting.

3. The third part of the document is a list of the names of the persons who were present at the meeting.

D'YAKONOV, I.A.; GOLODNIKOV, G.V.; REPINSKAYA, I.B.

Reactions of aliphatic diazo compounds with unsaturated compounds.
Part 25: Reaction of diphenyldiazomethane with silicon olefins.
Zhur.org.khim. 1 no.2:220-225 F '65.

(MIRA 18:4)

1. Leningradskiy gosudarstvennyy universitet.

D'YAKONOV, I.A.; REPINSKAYA, I.B.; GOLODNIKOV, G.V.

Trimethylsilylcarbene, a new methylene radical. *Izv. Akad. Nauk SSSR*.
35 no.1:199 Ja '65. [RUSSIAN]

L. Leningradskiy gosudarstvennyy universitet.

BOYAROV, I.A.; GOLODNIKOV, G.I.; PRIZHVA, L.F.

Reactions of aliphatic diazo compounds with unsaturated compounds.
Part 15: Reaction of ethyl ester of diazoacetic acid with
triisobutyl-, triisobutylallyl-, and triisobutyl- γ -benzylidene- α -methyl- β -methyl- γ -butyrolidone.
Zhur.b.khin. 35 no.12: 2181-2185, 1965.

(R11:10:1)

L. Leningradskiy Gosudarstvennyy universitet, Leningrad, 1964.

L 45894-60 ENT(m)/ENT(f) NM/RM

ACC NR: AP6026430

(A)

SOURCE CODE: UR/0079/56/036/005/0949/0949

AUTHOR: D'yakov, I. A.; Repinskaya, I. B.; Golodnikov, G. V.

ORG: Leningrad State University (Leningradskiy gosudarstvennyy universitet)

TITLE: Relative rate of addition of diphenylmethylene to trimethylvinyl-, trimethylallyl- and trimethyl- γ -butenylsilanes and 1-heptene

SOURCE: Zhurnal obshchey khimii, v. 36, no. 5, 1966, 49

TOPIC TAGS: silane, heptene, organosilicon compound

ABSTRACT: Diphenylmethylene (DM), obtained by thermal decomposition of diphenyldiazomethane (DD), adds to the silicoolefins (SO) trimethylvinyl-, trimethylallyl- and trimethyl- γ -butenylsilanes, forming silicon-containing cyclopropane (CD). An estimate of the relative activity of SO in their reaction with DD was made, based on the use of the method of competing reactions, and on the determination of the relative rate constant of the reaction (K_{rel}) of each of the SO. The reference standard was 1-heptene, whose rate constant of the reaction with DD was taken as unity. The K_{rel} values obtained were: $(CH_3)_3SiCH=CH_2$, 46.3; $(CH_3)_3SiCH_2CH=CH_2$, 0.93; $(CH_3)_3SiCH_2CH_2CH=CH_2$, 0.91. These values indicate that the steric effect of the trimethylsilyl group in trimethylvinylsilane has no appreciable effect on the rate of addition of DM to the double bond of this silane. The product of the addition of DM to 1-heptene (16% yield).

Card 1/2

UDC: 547.6+547.245

L 45894-66

ACC NR: AP6026430

1-amyl-2,2-diphenylcyclopropane, is described for the first time.

SUB CODE: 07/ SUBM DATE: 12Oct65/ ORIG REF: 003/ OTH REF: 004

Card 2/2 *LC*

65

GOLODNIKOV, N.A., master

Electric tachometer. *Energetik* 8 no.8:27-28 Ag '60. (MIRA 13:10)
(Tachometer)

GOLODNOV, A.; GUSEVA, N., red.; MAGIBIN, P., tekhn. red.

[Sheep raising as a basic source of meat production]
Ovtsevodstvo - osnovnoi istochnik proizvodstva miasa.
Alma-Ata, Kazsel'khozgiz, 1962. 26 nos. in 1 v. 22 p.
(MIRA 17:1)

SALYUKOV, P.A., kand. biol. nauk, VERNIGOR, V.A., kand. sel'khoz. nauk; KOR'ANOVSKAYA, M.A., kand. sel'khoz. nauk; GOLOLNOV, A.V.; SKOROBOGATOV, Yu.A., mladshiy nauchnyy sotr.; MALLITSKIY, V.A., kand. sel'khoz. nauk; CHAPCHIN, B.V., kand. sel'khoz. nauk; PONOMAREV, P.P., kand. tekhn. nauk; BARVINTSEV, Yu.N., doktor sel'khoz. nauk; NECHAYEV, I.N., mlad. nauchnyy sotr.; POZDNYAKOV, P.M., kand. biol. nauk; KOVIN'KO, D.A., kand. biol. nauk; BAIANINA, G.V., kand. sel'khoz. nauk; MOISEYEV, K.V., kand. sel'khoz. nauk; ROMANOV, P.F., kand. veter. nauk; PAI'GOV, A.A., kand. veter. nauk; ANAN'YEV, P.K., kand. veter. nauk; VASIL'YEV, B.M., kand. sel'khoz. nauk; ABDULLIN, V.A., kand. ekon. nauk; GALIAKBEROV, N., laureat Gos. premii, kand. sel'khoz. nauk, red.; GUSEVA, N., red.; NAGIBIN, P., tekhn. red.

[Reference book for zootechnicians] Spravochnik zootekhnika.
Pod red. N. Galiafterova. Alma-Ata, Kazsel'khozgiz, 1963.
492 p. (MIRA 16:5)

(Kazakhstan--Stock and stockbreeding)

1990

"In addition, neither of the names listed in the caption of the exhibit, 'C. L. Taylor, Jr.' or 'John Henry George Taylor, Jr.', listed as 'John Elbertson'."

Disertationsschriftel der Medizinischen Fakultät der Universität
München, April 1967.

THE UNIVERSITY OF CHICAGO

AID P - 3435

Subject : USSR/Electricity
Card 1/2 Pub. 27 - 2/32
Author : Golodnov, M. N., Kand. of Tech. Sci., Dotsent,
Rostov-on-the-Don
Title : Permissible load in starting synchronous motors with
directly-connected exciter
Periodical : Elektrichestvo, 10, 7-10, 0 1955
Abstract : Current circulars of the Ministry of Electric Power
Stations No. 7/E 1952 and No. E-5/54 1954 permit the
starting without preliminary tests of synchronous
motors with directly connected exciter and with
resistance moment not exceeding 0.4 of the nominal.
The author presents a simple engineering method of
approximate calculations to determine the possibility
of applying direct starting of synchronous salient-
pole motors. He presents results of calculating the
characteristics of motors of the SM-300-750 217-kw,

Elektrichestvo, 10, 7-10, 0 1955

AID P - 3435

Card 2/2 Pub. 27 - 2/32

6-kv, and SM-160-500 113-kw, 380-v types. The author suggests experimenting with his method to ascertain the possibility of starting motors with a resistance moment above 0.4 of the nominal. Analytical findings of the author have been verified experimentally not only by himself, but also by several laboratories (Central Scientific Research Institute of the Ministry of Electric Power Stations and others). Two diagrams, 2 Soviet references (1950, 1952).

Institution : None

Submitted : N 29, 1954

AUTHORS: Golodnov, L. N., Docent, Candidate of Technical Sciences (Moscow) 105-98-4-22/37

TITLE: Experimental Determination of the Parameters of an Equivalent Generator in Linear A. C. Circuit
(Opytnoye opredeleniye parametrov ekvivalentnogo generatora v lineynykh tsepyakh peremennogo toka)

PERIODICAL: Elektrichestvo, 1958, Nr 4, pp. 77-78 (USSR)

ABSTRACT: The determination of the electromotive force of an equivalent generator can be determined without difficulty from the idle motion of a dipole. It is useful to determine the total resistance of the equivalent generator in trying to feed the dipole from a foreign source. This should be done by measuring it at the terminals of the dipole as input resistance while all energy sources feeding it are disconnected and replaced by circuit sections with the same internal resistances. Sometimes, however, such a disconnection is not possible or is not desired. In such a case the parameters r_H and x_H can be determined by two experiments applying different loads to the effective dipole.

Card 1, 3

Experimental Determination of the Parameters of an
Generator in Linear A. C. Circuit

105-58-4-22/57

After the idle motion voltage $U_0 = E_H$ was determined the load resistance $Z_1 = r_1 + jx_1$ is connected to the dipole terminals. Then

$$I_1 = \frac{U_0}{\sqrt{(r_H + r_1)^2 + (x_H + x_1)^2}} \quad (1)$$

I_1 is determined according to the amperemeter indication and r_1 and x_1 , if unknown, by the indications of amperemeter, voltmeter and wattmeter. The resistance Z_1 is replaced by $Z_2 = r_2 + jx_2$ and the equation (2) is obtained. Both equations (1) and (2) are solved in common and the resistance r_H and x_H are determined. In practice mostly only effective resistances r_1 and r_2 can be connected. Thus

Para 2/3

Experimental Determination of the Parameters of an
Equivalent Generator in Linear A. C. Circuit

105-58-4-2/37

the scheme of the experiment becomes more simple (the
wattmeter is no longer needed). The method was tried in
practice and the results were good.

AVAILABLE: Library of Congress

1. Generators-Parameter determination-Theory 2. Linear circuits

Card 3/3

AUTHOR: Golodnov, M. N., Candidate of Technical Sciences 1c5-56-6-20, 11
(Rostov-na-Donu = Rostov on Don)

TITLE: The Determination of the Synchronous Reactance of Three-
-Phase-Current Generators With Salient Poles and Permanent
Magnets in an Experimental Way (Opredeleniye opytnym putem
sinkhronnogo reaktivnogo soprotivleniya trekhfaznykh yavno-
polyusnykh generatorov s postoyannymi magnitami)

PERIODICAL: Elektrichestvo, 1958, Nr 6, pp. 91-91 (USSR)

ABSTRACT: The method of the small slip which is employed for the deter-
mination of the synchronous reactances x_d and x_q of motors
with salient poles cannot be used for motors with permanent
magnets. In this case it is expedient to employ an indirect
method for the determination of x_d and x_q by means of a test
with two loads. This method is described here. During idling
of the generator the linear voltages U_{ab} , U_{bc} , U_{ca} , are mea-
sured and the no-load e.m.f. E_0 is determined ... equation(1).
An inductive symmetrical three-phase load is connected with
the generator, where the load possesses a low effective re-
sistance. ($\cos \phi = 0.1-0.2$). Then the amperages and the linear
voltages of the generator are measured. Corresponding to the

Card 1/2

The Determination of the Synchronous Reactance of Three-Phase-Current Generators With Salient Poles and Permanent Magnets in an Experimental Way

vector diagram the formula (2) is then written down and from it the formula (3) for x_d is obtained. As the angle φ is near to 90° and differs little from Ψ , (3) can be written down in the form of (4). For the determination of x_q a symmetrical load of a capacity nature is produced in a way that current I according to its phase coincides with the e.m.f. E_0 . x_q is determined from equation (6). There are 2 figures.

1. Generators--Performance 2. Generators--Analysis 3. Magnets
--Electrical effect --Mathematics

Card 2/2

AUTHOR: Golodnov, Mikhail Nikolayevich, Candidate of Technical Sciences, Docent SOV/144-58-7-5/15

TITLE: Equations of the Additional Current and Torque that Characterise the Influence of the Exciter e.m.f. on Asynchronous Operation of a Synchronous Machine (Uravneniya dopolnitel'nogo toka i dopolnitel'nogo momenta, kharakterizuyushchiye vliyaniye e.d.s. vzbuditelya na asinkhronnyuyu rabotu sinkhronnoy mashiny)

PERIODICAL: Izvestiya vysshikh uchebnykh zavedaniy, Elektromekhanika, 1958, Nr 7, pp 57-66 (USSR)

ABSTRACT: The meaning of the term 'additional currents' is first defined as follows: If the equations of a synchronous machine are expressed in operator form the stator and rotor currents may be divided into two components. The first of these are currents due to the system voltage applied to the stator circuit which do not depend on the exciter e.m.f. These currents can be observed under asynchronous (or synchronous) conditions in the absence of excitation and they are termed 'asynchronous currents'. Secondly, the currents due to the presence of the exciter e.m.f. in the closed field circuit. In

Card 1/6

SOV/144-58-7-6/15

Equations of the Additional Current and Torque that Characterise the Influence of the Exciter e.m.f. on Asynchronous Operation of a Synchronous Machine

the presence of excitation these currents are superposed on the 'asynchronous currents' and are termed 'additional' currents. If the effects of saturation and hysteresis are neglected the effect of the asynchronous and additional currents may be considered separately. The equations for the additional current and the additional torque together with the equations for the asynchronous current and torque may be used to analyse the operation of synchronous machines running under asynchronous conditions with the exciter connected. In particular these equations have been useful in analysis of the following conditions: starting of synchronous motors with the exciter connected; self-synchronisation of generators; asynchronous operation when the field circuit is open; and some cases of loss of synchronism of a motor. The operator equations for the additional currents are given in expressions (10) - (16). Expressions are then derived for the additional current when the field

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current is constant, which is the case for all practical purposes if the exciter is left connected and the field rheostat is not suddenly altered during the process. A number of practical conclusions are derived from the equations. Test results of the additional current and short circuit current as functions of the field current are given in Fig 2 for nine different motors; the test results are stated to confirm equations (31) - (33) and the conclusions drawn from them. The additional current was determined from oscillograms as shown in Fig 1. The characteristics obtained were compared with short circuit characteristics and were usually found to coincide, but in some cases the additional current characteristic lies somewhat lower than the short circuit characteristic, though as will be seen from Fig 2 the difference is not great. The torque expression (3) contains some terms which depend on the angle δ and some which do not. Accordingly the additional torque can be resolved into two components, a mean value given by expression (34)

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and a periodic component given by expression (35). By further analysis it is shown that the periodic component of the additional torque does not depend on the slip and varies sinusoidally with the slip frequency. An oscillogram of this torque as a function of δ with constant slip is given in Fig 3. Expression (39) is derived for the mean value of the additional torque. It will be seen that in general the mean additional torque is small and it is important only in a few special cases. One case in which the additional torque is important is when on starting a synchronous motor against a load the machine fails to run up to synchronous speed because of troughs in the torque/speed curve. It is shown that in some cases if the field current is increased when this happens the torque may become less because of the retarding effect of the mean additional torque, and then it is necessary either to reduce the mechanical load or to introduce active resistance into the field circuit so that the machine can run up to

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speed. Test data showing results of increasing the field
current when machines fail to run up to speed are given
in Table 1, which also includes a few torque values.
The assumptions made in the article limit the field of
application of eqs (29), (33), (38) and (39) to
conditions where the slip is much less than 1, in large
modern machines of fairly large flywheel effect with low
active stator resistance and a massive rotor. The
limitations imposed in the derivation of formulae (38)
and (39) render them unsuitable for use at half
synchronous speed. Thus Eqs (29) and (38) and
conclusions drawn from them can be used for the
relatively common case of a machine operating under
synchronous conditions with low active resistance and
slip less than 0.5. Under other conditions and
particularly at low speeds there may be other components
of additional current and torque which may be of

Card 5/6

SOV/1944-72-7-4/15

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considerable importance. In such cases it is best to
use Eqs (20) - (24) or graphical methods which are not
considered here.

There are 3 figures, 1 table and 9 Soviet references.

ASSOCIATION: Kafedra elektrotekhniki Rostovskogo instituta
sel'skokhozyaystvennogo mashinostroyeniya (Chair of
Electrical Engineering Rostov Institute of
Agricultural Machine Building)

SUBMITTED: January 31, 1956

Card 6/6

GOLODNOV, Yu.M.

New method for determining the angle between electromotive force
vectors and the voltage on the rings of the transformer.
Sbor.rats.predl.vnedr.v proizv. no.5:50-52 '60. (MIRA 14:8)

1. "Yuvenergochermet".
(Electric current converters)

GOLODNOV, Yu.M., inzh.

Some characteristics of a rectifier cascade. Vest.elektrom.
31 no.1:21-24 Ja '60. (MIRA 13:5)
(Electric motors, Induction) (Electric driving)

9.2200 (1001,1482)

3219h
S/196/61/000/010/022/037
E194/E155

AUTHOR: Golodnov, Yu. M.

TITLE: A direct current amperes-squared-hour meter

PERIODICAL: Referativnyy zhurnal, Elektrotehnika i energetika,
no. 10, 1961, 1 abstract LOK 6 (Vestn elektroprom-
sti, no. 4 1961, 56-60)

TEXT: "Yuvenergochermet" has developed and made a direct
current amperes-squared-hour meter, using compact amplidyne.
The electrical circuit and winding data are given, and the
construction and operating principles are described. The meter
reading is completely independent of the direction of current in
the amplidyne control winding. Any single-phase induction meter
for 127 V, 5 A can be used in the circuit. With a rated control
winding current of 2.5 A the meter constant is 6.52 A²h/kWh. The
equipment fits into a carrying case and is easily transported. ✓
A variant is described in which the secondary winding of the
amplidyne carries 5 A. This set consumes less power, so that a
150 VA stabiliser can be used. With a rated control winding
current of 1.5 A the meter constant is 0.965 A²h/kWh.

Card 1/2

32191

A direct current amperes-squared.

S/196/61/000/010/022/037
E194/E155

An important advantage of the meter is its ability to operate with a shunt for 33 - 45 mV from which it draws a very small current. Shunts for 75 and 100 mV are used in conjunction with additional resistances. The meter can be used to measure r.m.s currents in reversing circuits having peak loads with an accuracy of 2.5%. The meter can be used in circuits of 100 A and more 5 figures

X

[Abstractor's note Complete translation]

Card 2/2

GOLODNOV, u.m., inzh.

D.C. ampere square hour meter. Vest. elektropram. 32 no.4:56.60
Ap 161. (MIRA 15:5)

(Electric meters)

(Electric machinery--Measurements)

SATPATAVA, T. I.; GOLUB, V. A.; N. I.

Isopyrite in the Dzhezkazgan mines. (Izv. V. N. Karadzh. Ser. geol.
1977, 10: 112-117, (MLBA 10:8)
(Dzhezkazgan - Soviet Union).

GOLODNOVA, O.S., inzh.

Review of L. A. Mirenburg's book "Repair of turbogenerator rotors
under stationary conditions." Elek. sta. 31 no.12:88 D '60.

(MIRA 14:5)

(Turbogenerators—Maintenance and repair)
(Mirenburg, L.A.)

GURVICH, V.S., inzh.; GOLODNOVA, O.S., inzh.

Rotor bands from aluminum alloys. Elek.sta. 32 no.4:94-95 Ap
'61. (MIRA 14:7)

(Turbogenerators)

GOLODNOVA, O.S., inzh.; DEGIL', G.S., inzh.; PANCHENKO, A.U., inzh.;
TUROS, A.E., inzh.; MESHKOV, V.X., inzh.

Concerning the seals of hydrogen cooled turbogenerators. Elek.
sta. 33 no.8:60-68 Ag '62. (MIRA 15:8)

1. Rostovenergo (for Golodnova). 2. Glavnoye upravleniye
energeticheskogo khozyaystva Donetskogo basseyna (for Degil',
Panchenko, Turós). 3. Moskovskoye rayonnoye upravleniye
energeticheskogo khozyaystva Glavtsentroenergo Ministerstva
elektrostantsiy SSSR (for Meshkov).
(Turbogenerators)

GOLODNOVA, O.S., inzh.

Choice of the dimensions of shaft grease seals. Vest. elektrom.
33 no.8:72-73 Ag '62 . (MIRA 15:7)
(Turbogenerators—Equipment and supplies)

GOLODNOVA, O.S., inzh.

Expenditure of hydrogen in the cooling of turbogenerators. Elek. sta.
36 no.6, 50, 53 Je '65. (MIRA 1887)

GOLODNYAK, G. S.

Automation of production processes at the Semipalatinsk cement
plant. TSement 27 no.5 24-27 S-O '61. (MIRA 14 13)

1. Semipalatinskiy tsementnyy zavod.
(Semipalatinsk - Cement plants)

GOLODNYAK, G.S.

Automatic sprinkling of the shells of cement mills. Tsement 28
no.1:21 Ja-F '62. (MIRA 16:5)

1. Bezmeinskiy tsementnyy zavod.
(Milling machinery)

GOLODNYAK, N.

Encourage the activities of key personnel. Voen.znan. 35 no.4:
11-12 Ap '59. (AIRA 12:7)

1. Predsedatel' Kiliyskogo rayonnogo Komiteta Vsesoyuznogo
dobrovol'nogo obshchestva sodeystviya armii, aviatsii i flotu.
(Military education)

USSR / Farm Animals. Small Horned Sheep

Abstract: Ref Zhur-Biol., No 5, 1958, 21471

Author : Olenin F. S., Gritsay S. P., Golodnyy I. F.

Inst :

Title : A New Technique of Feeding Lambs and Its Effectiveness (Novaya tekhnika kormleniya yagnyat i ego ekonomicheskaya effektivnost')

Orig Pub: S. kh. Povolzh'ya, 1957, No 6, 67-69

Abstract: Laboratory experiments were carried out in order to test "rhythmic" feeding onweanling lambs (the ration of one 6-day period was increased by 20% and that of the other 6-day period was diminished by 20%). It was found that the average daily weight increase of test animals was 32%, shearing yield 400 g. more, and the feeding per 1 kg. of weight increase was 2.42 feed units less than in the control group.

Card 1/2

AUTHOR: *Golodnyy, Ts.* 25-11-24/23
 TITLE: Explorer of Arctic Regions (Issledovatel' Arktiki)
 PERIODICAL: Nauka i Zhizn', 1957, # 11, pp 57-58 (USSR)
 ABSTRACT: Petr Filimonovich Shvetsov, doctor of geological-mineralogical sciences, member-correspondent of the USSR Academy of Sciences, winner of the Stalin prize, succeeded the Soviet geologist Vladimir A. Obruchev as head of the Institute for the Study of Congelation. Soviet scientists, such as Gumgin, Tsytovich, Tolstikhin, Kudryavtsev, Baranov, Soltykov, prepared the ground for this new branch of science. Shvetsov was among the first scientists to recommend the exploration of the arctic regions. He established the first Soviet station for the study of congelation on the Chukotsk peninsula. His first scientific publication dealt with the research of the Pinksheynsk thermal spring, and he provided the possibility to take advantage of these valuable resources. The results of his second expedition to the Chukotsk peninsula were summed up in his second work "Eternal Congelation and Technical Geological Conditions in the Anadyrsk Region". The material for his latest publication

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Explorer of Arctic Regions

25-11-24/28

"Subterranean Waters of the Verkhoyansk-Kolyma Folded Region and Peculiarities of Their Occurrence, Connected with Eternal Congelation", was collected during a two-year research expedition to Yakutiya.

There are three photographs and one sketch.

AVAILABLE: Library of Congress

Card 2/2

GOLODNYI, TS .

~~Production line~~ system in building apartment houses. Nauka
i zhizn' 24 no.3:21-24 Mr '57. (MLRA 10:5)
(Apartment houses)

62-100000-13
AUTHOR: Golodnyy, Ts.

86-1-2004

TITLE: An Experimental Block (Kvartal. eksperimentalny)

PERIODICAL: Nauka i Zhizn', 1966, # 1, pp 60-64 (USSR)

ABSTRACT: This article deals with the construction of the first Soviet experimental apartment buildings in the Novyye Cheremuski area of Moscow. The building ground consists of two parts: the housing area and the social center. There are 13 four-storied and 5 eight-storied apartment houses under construction. The social center consists of a school, a kindergarten, a public nursery, repair services, shops, a public dining-room and parking lots.
There are seven photographs and one illustration.

AVAILABLE: Library of Congress

Card 1/1

GOLODNYI, TS.

At the anniversary exhibition. Rabotnitsa 36 no.1:24 Ja '58.
(MIRA 11:2)
(Moscow--Art--Exhibitions)

GOLODNYI, TS.

A book arrived at the plant. Rabotnitsa 37 no.10:27 0 '59.
(MIRA 13:2)

1. Kombinat "Trekhgornaya manufaktura," Moskva.
(Moscow--Booksellers and bookselling)